

LARIONOV, H.I.

Automatizing the machining of bearing rings. Avt.i trakt.prom.
no.8:46 Ag '57.

1. Mauchno-issledovatel'skiy institut transports avtomobil'noy
promyshlennosti.

(Bearing industry)

28(1) SOV/118-59-4-4/25

AUTHORS: Lysyakov, A.G., Engineer; Preobrazhenskiy, M.A.,

Candidate of Technical Sciences; and Larionov, N.I.,

Engineer

TITLE: Bridge-Type Stacking Cranes

PERIODICAL: Mekhanizatsiya i avtomatizatsiya proizvodstva, 1959,

Nr 4, pp 14-17 (USSR)

ABSTRACT: The design office of the Nauchno-issledovatel'skiy

institut tekhnologii avtomobil'noy promyshlennosti (Scientific Research Institute of Technology of the Automobile Industry) has developed, under the supervision and with the participation of the Vsesoyuznyy nauchno-issledovatel'skiy institut pod"ëmnogo-transportnogo mashinostroyeniya (All-Union Scientific Research Institute of Lifting and Transportation Machine Building), a bridge-type stacking crane for the semifinished product warehouse of the Moskovskiy zavod

malolitrazhnykh avtomobiley (the Moscow Small

Card 1/2 Car Plant). Technical characteristics are:

ARKHIPOV, N.A.; LARIONOV, N.I.

Automatic machine for the manufacture of AKF-2 shell molds.

Avt.prom. 27 no.6:46 Je '61. (MIRA 14:6)

l. Nauchno-issledovatel'skiy tekhnologicheskiy institut avtomobil'noy promyshlennosti.

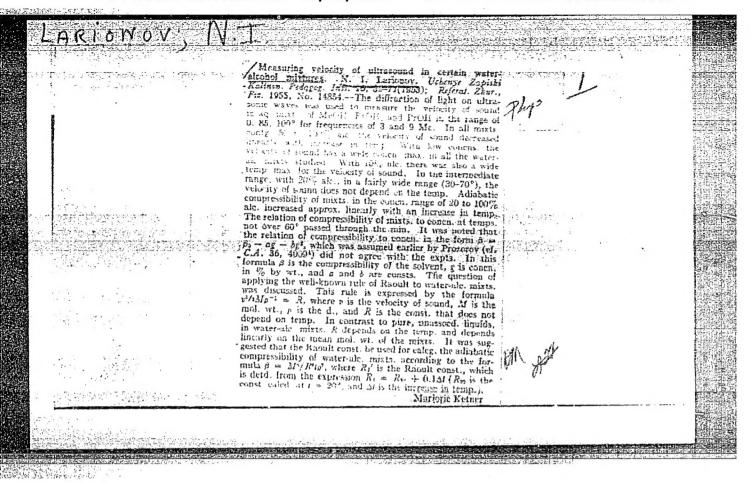
(Automobile industry--Equipment and supplies)

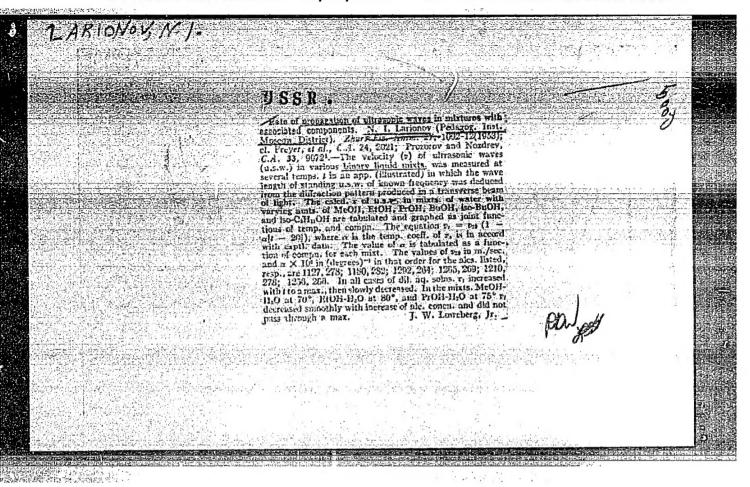
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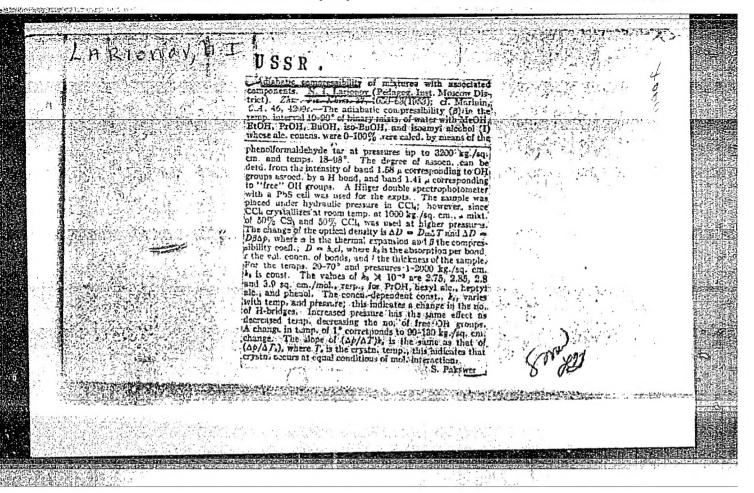
"Investigation of Adiabatic Congressibility of Fintures With Associated Comments." Sub 25 Dec 51, Moscow Chlast Pedagogical Inst.

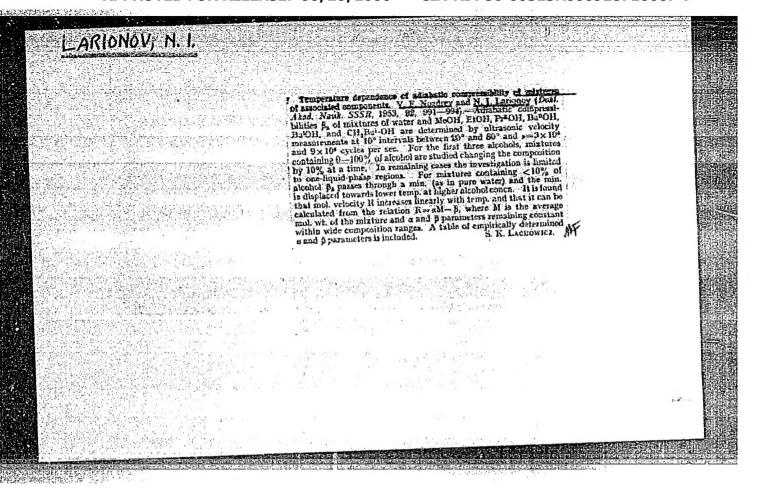
Discertations presented for science and entineering degrees in Possess during 1911.

SO: Sum. No. 480, 9 May 55.









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CGRYACHKO, C. V., DMITRIYEVA, NA. end LARIOMOV, N. I.

"Acceleration of the Dyeing of Synthetic Mbers."

report presented at the 6th Sci. Comfgrence on the Application of Ultrasound in the investigation of Matter, 3-7 F b 1956, organized by Min. of Education RSFER and Moscow Oblast Pedagogic Inst. im N. K. Krupskaya.

GORYACHKO, G.V.; LARIOMOV, N.I.; GIAZKOVSKIY, Yu.V.

Ultrasonic cleaning of spinnerets. Khim.volok. no.1:51-52
'60. (MIRA 13:6)

1. Kalininskiy pedinstitut (for Goryachko, Larionov). 2. Kalininskiy filial Vsesoyuznogo nauchno-issledovatel'skogo instituta iskusstvennogo volokna (for Glazkovskiy).

(Bayon spinning)

(Ultrasonic waves—Industrial applications)

24.1800

65961

SOV/58-59-4-9090

Translation from: Referativnyy Zhurnal Fizika, 1959, Nr 4, p 245 (USSR)

AUTHORS:

Larionov, N.I., Dmitriyeva, N.A., Goryachko, G.V.

TITLE:

Study of the Physical and Chemical Properties of Aqueous Solutions of Dimethylformamide\by the Ultrasonic and Other Methods in the 20° to

PERIODICAL:

V sb.: Primeneniye ul'traakust. k issled. veshchestva, Nr 7, Moscow, 1958, pp 75 - 90

ABSTRACT:

The authors submit the results of measuring the velocity of propagation and the molar velocity of ultrasonic waves, as well as the density, adiabatic compressibility, viscosity, surface tension, and refractive index in aqueous solutions of dimethylformamide (D) at concentrations ranging from 0% to 100% (at 10% intervals) and at various temperatures. The data are presented in the form of tables and curves. At 20°C the ultrasonic velocity passes through a maximum at a concentration of 50 wt %, the density up to a concentration of 60 wt.% is close to that of pure water, the adiabatic compressibility of D solutions passes through a minimum at a concentration of 50 wt.%, the viscosity shows

Card 1/2

65961

SOV/58-59-4-9790

Study of the Physical and Chemical Properties of Aqueous Solutions of Dimethylformamide by the Ultrasonic and Other Methods in the 20° to 90°C Range

a well-defined maximum at a concentration of 60 wt.%, the refractive index rises monotonously with the concentration, and the surface tension drops monotonously. The molar ${\cal A}$ velocity of sound in aqueous solutions of D increases with an increase in the temperature and concentration. (Kalininskiy ped. in-t).

A.A. Senkevich

Card 2/2

APPROVED FOR RELEASE: 06/20/2000 CIA-RDP86-00513R000928710007-7"

SOV/58-59-5-11512

Translation from: Referativnyy Zhurnal Fizika, 1959, Nr 5, p 228 (USSR)

Goryachko, G.V., Dmitriyeva, N.A., Larionov, N.I. AUTHORS:

Use of <u>Ultrasonic Waves</u> to Dye Polyacrylonitrile "Nitron" Fiber TITLE:

V sb.: Primeneniye ul traakust. k issled. veshchestva. Nr 7, Moscow, PERIODICAL:

1958, pp 161 - 167

The authors report on the results of using ultrasonic waves (US) to dye polyacrylonitrile "nitron" fiber. It was established that the dry ABSTRACT: fiber is rapidly and permanently dyed with the aid of the dispersed dyes for acetate silk by a method involving the combined use of US and accelerants. The preliminary dispersing of the dye by means of US tells favorably on the dyeing rate and the utilization of the dye; in this connection it is more advantageous to use lower frequencies. Using US in conjunction with an accelerant (aniline) cuts the dyeing time in half

as compared with the case of using the accelerant alone. The emulsions prepared under the influence of US yield better results. (Kalininskiy ped, in-t, USSR).

A.A. Senkevich Card 1/1

GELLER, B.E.; GORYACHKO, G.V.; DMITRIYEVA, N.A.; LARIOMOV, N.I.

Destruction of polyacrylonitrile by the action of an ultrasonic field. Vysokom.soed. 1 no.11:1610-1616 H '59. (MIRA 13:5)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut iskusstvennogo volokna i Kalininskiy pedagogicheskiy institut.

(Acrylonitrile) (Ultrasonic waves)

LARIONOV, N.I.

PHASE I BOOK EXPLOITATION SOV/5644

Vserossiyskaya konferentsiya professorov i prepodavateley pedagogicheskikh institutov

Primeneniye ul' traakustiki k issledovaniyu veshchestva. vyp. 10. (Utilization of Ultrasonics for the Investigation of Materials. no. 10) Moscow, Izd-vo MOPI, 1960. 321 p. 1000 copies printed.

Eds.: V. F. Nozdrev, Professor, and B. B. Kudryavtsev, Professor.

PURPOSE: This book is intended for physicists and engineers interested in ultrasonic engineering.

COVERAGE: The collection of articles reviews present-day research in the application of ultrasound in medicine, chemistry, physics, metallurgy, ceramics, petroleum and mining engineering, defectoscopy, and other fields. No personalities are mentioned. References accompany individual articles.

Card 1/10

Utilization of Ultrasonics (Cont.) SOV/5644 TABLE OF CONTENTS: Nozdrev, V. F. Physical Principles of the Engineering and Technical Use of Low-Amplitude Molecular Acoustics Larionov, N. I., G. V. Goryachko, N. A. Dmitriyeva, and B. E. Geller [Kalininsk. pedinstitut im. M. I. Kalinina, Kalininsk. filial VNIIV-Kalinin Pedagogical Institute imeni M. I. Kalinin, Kalinin Branch of the All-Union Scientific Research Institute for High Polymers]. Investigation of Degradation Processes in High Polymers Under the Action of an Ultrasonic Field 23 Kogan, I. N., L. I. Menes, and N. I. Parlashkevich [N.-i. int plastmass - Scientific Research Institute for Plastics]. Continuous Measurement of Viscosity With the Aid of an Ultrasonic Viscometer Card 2/10 33

26254 S/194/61/000/001/021/038 D216/D304

15.8530

Larionov, N.I., Goryachko. G.V., Dmitriyeva, N.A. AUTHORS:

and Geller, B.E.

Analysis of the high polymer degradation process TITLE:

under the influence of an ultrasonic field

Referativnyy zhurnal. Avtomatika i radioelektronika, no. 1, 1961, 15, abstract 1 El31 (V Sb. Primeneniye ul'traakust. k issled. veshchestva, no. 10, M., PERIODICAL:

1960, 23-32)

The results are given of experimental analysis of the process of degradation of polyacrylonitrile (N/H (PAN)) and of other forms of polymers (e.g. acetyl cellulose AU - (ATs) in the solution of dimethylformamide $\Box M\Phi$ (DMF)) under the action of a powerful ultrasonic field as a function of frequency and power for concentrations up to 5 g/l. It is shown that under the action of a field intensity up to 20 W/cm² and frequency 500 Kc/s, the molecules of

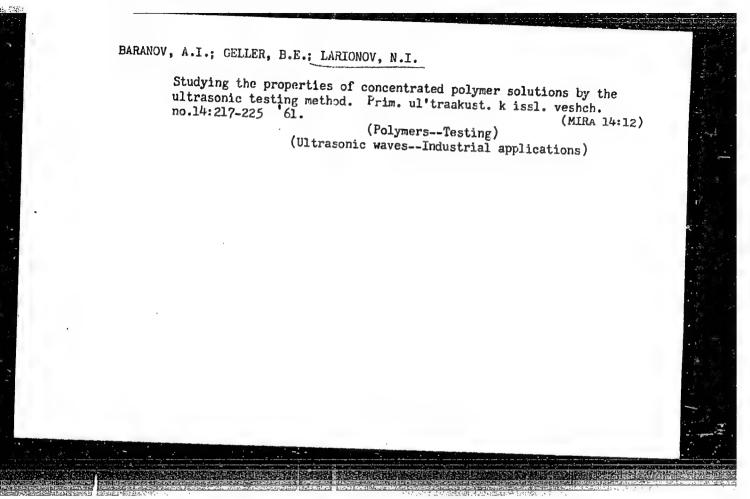
Card 1/2

25254 S/194/61/000/001/021/038 D216/D304

Analysis of the high polymer...

PAN are degraded while those of acetylcellulose remain unchanged. This fact is explained by the strength of chemical bonds within the polymer structure between the polymer molecule and the side-groups. The kinetics of the degradation process of PAN have been studied. It is shown that the depolymerization process follows the lst order reaction and that long chain molecules are degraded first. The results are shown in the form of graphs. 22 references.

Card 2/2



3513h \$/058/62/000/002/019/053 A058/A101

1.1800

AUTHORS: Tsepelev, A. I., Larionov, N. I., Mikhaylov, F. G.

TITLE: Ultrasonic effect in the process of galvanic plating

PERIODICAL: Referativnyy zhurnal, Fizika, no. 2, 1962, 43, abstract 2G321

(V sb. "Primeneniye ul'traakust. k issled. veshchestva", no. 14,

Moscow, 1961, 227-230)

TEXT: Nickel-plating in an ultrasonic field enables one to increase the current density 2-3 times over and to carry out nickel-plating at reduced temperatures (20 - 30°C); at the same time, the quality of the nickel platings is improved. The isotherms of current density as a function of ultrasonic power were found. It was found that the highest yield of chromium per current takes place when ultrasonic intensity = 1 watt/cm².

[Abstracter's note: Complete translation]

Card 1/1

1,1700

AUTHORS:

Tsepelev, A. I., Larionov, N. I. and Mikhaylov, F. G.

TITLE: .

The influence of ultrasound on the galvanic-coating

process

PERIODICAL:

Referativnyy zhurnal, Avtomatika i radioelektronika, no. 4, 1962, abstract 4-5-40g (V sb. Primeneniye ul'traakust. k issled. veshchestva. no. 14, N., 1961,

227-230)

TEXT: It is established that ultrasound of 22 kc/s enables one to increase the current density by 2 - 3 times and to carry out the nickel-plating process at a lowered temperature. The optimum ultrasonic intensity for the largest output of chromium for a given current is determined. The magnetostrictor is so placed that ultrasound propagates parallel to the surface of the object. The process of degreasing of the object before coating was intensified by ultrasound. 2 references. / Abstracter's note: Complete translation. /

Card 1/1

S/275/63/000/001/030/035 D413/D308

Larionov, N. I. and Goryachko, G. V. AUTHORS:

The degradation of acetyl-cellulose under the action TITLE:

of an ultrasonic field

Referativnyy zhurnal, Elektronika i yeyo primeneniye, PERIODICAL:

no. 1, 1963, 17, abstract 1V 123 (In collection: Primenentye ul traakust. k issled. veshchestva, no. 15,

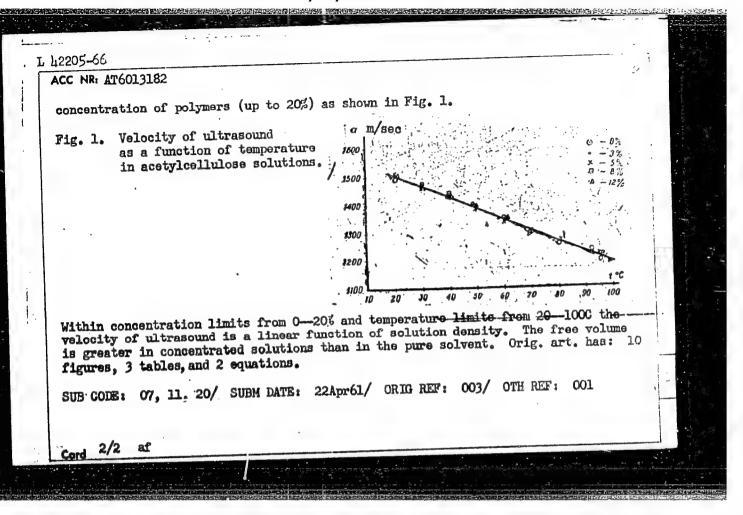
M., 1961, 249-254)

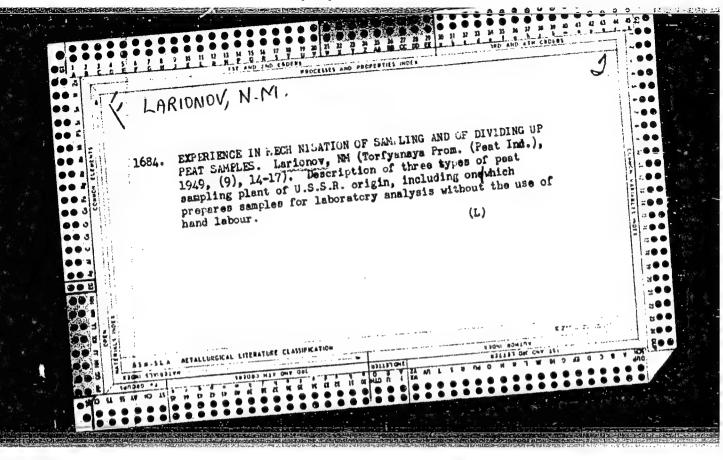
TEXT: The authors have investigated the degradation of acetyl cellulose, dissolved in dimethylformamide (DMF), under the action of 300 kc/s ultrasonic vibration at an intensity of up to 20 W/cm². The range of concentrations 1 - 5 g/liter acetyl cellulose in DMF was used in this work, since in more concentrated solutions the results may be affected by factors related to the interaction of macromolecules. It was shown that under the action of an intense ultrasonic field the viscosity of solutions of acetyl cellulose in DMF decreases with time. A minimum viscosity is maintained for a

Card 1/2

The degradati	m of	S/275/63/000/001/03 D413/D308	0/035
not to disagg macromolecule solutions of degradation o cates the eff	regation of the solutions of acetyl cellulose acetyl cellulose in D solutions of polyacect of the strength of the serion of an interest of the serion of an interest.	that the fall in viscosity ion but to degradation of t. The process of degradation MF proceeds more slowly that rylonitrile in DMFm which if the chemical bonds in the ense ultrasonic field, while the chemical to a change of the change to a change the state of the change to a change the state of the change the c	n of n the ndi- poly- e in
causing degra	wave velocity in th	, does not lead to a change e solutions. 8 references.	Ab
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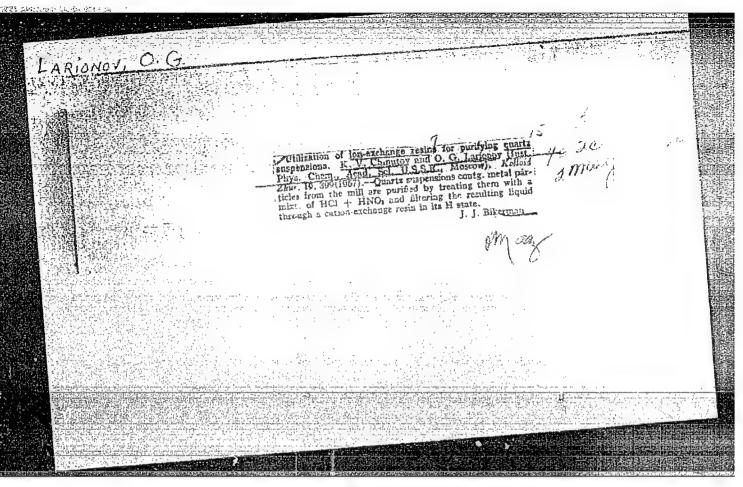
I 42205-66 EWT(m)/EWP(j)/T/EWP(k) ACC NR: AT6013182 SOURCE CODE: UR/0000/61/000/600/6017/02 AUTHORS: Baranov, A. I.; Geller, B. E.; Larienov, N. I. ORG: none TITLE: Study of the properties of concentrated polymer solutions using an ultrasound method SOURCE: Moscow. Oblastnoy pedagogicheskiy institut. Primeneniye ul'traakustiki k issledovaniyu veshchestva, no. 14, 1961, 217-225 TOPIC TAGS: ultrasound, ultrasonic velocity, fluid density measurement, salution Aconcentration, adiabatic compression, ultrasonic wave propagation ABSTRACT: The relationship between the density, temperature, and concentration of goncentrated solutions of perchlorovinyl, polyacrylonitrile, and acetylcelluloge was investigated using ultrasonic methods. The work is presented as a part of a complex effort by N. I. Larienev, G. V. Gorvachko, N. A. Dmitriyeva, B. E. Geller (\$b. Primeneniye ul'traakustiki k issledovaniyu veshchestva, vyp. X, str. 23, H., 1960), designed to study physical and chemical properties of polymers. Dimothylformamide was selected as the solvent. Changes in density, in the propagation velocity of ultrasound, and in adiabatic compressibility of the solutions were measured. The velocity of ultrasound was measured optically, with an accuracy of 1%, It was established that the propagation velocity is practically independent of the Card 1/2

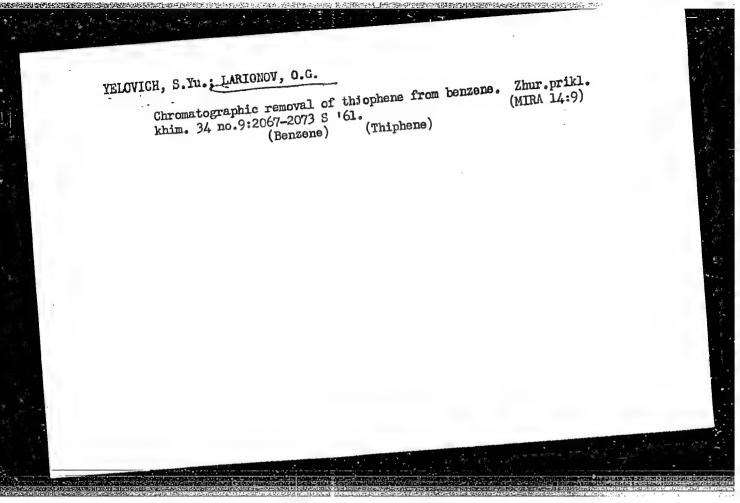


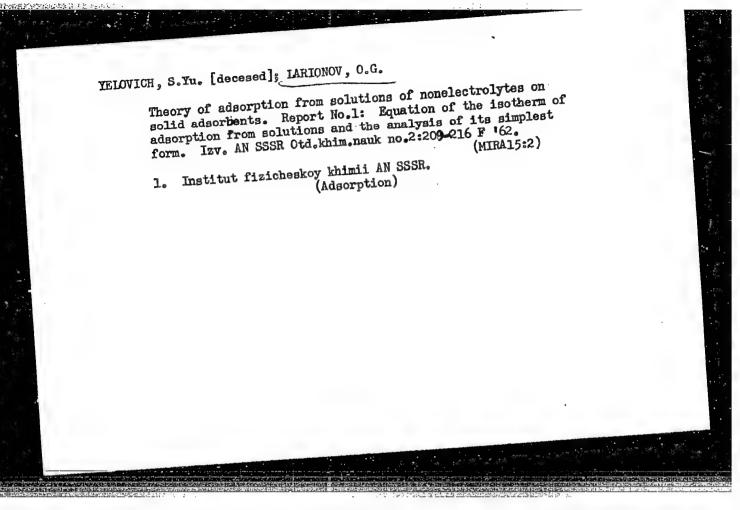


- 1. SKYUYEV, P.V., LARIONOV, N. V., Eng., SAPRYGIN, I. S.
- 2. USSR (600)
- 4. Metals Head Treatment
- 7. Reducing stresses in parts by annealing. No 9 1952.

9. Monthly List of Russian Accessions, Library of Congress, April 1953, Uncl.







YEIOVICH, S.Yu. [deceased]; LARIONOV, O.G.

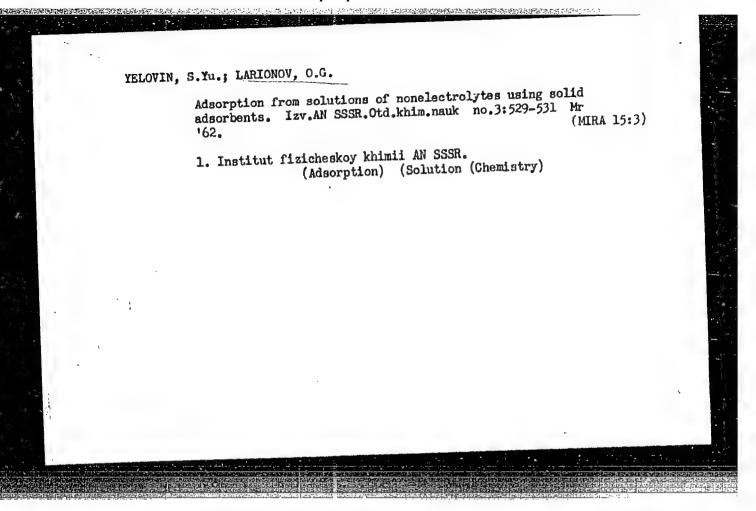
Theory of adsorption from solutions of nonelectrolytes on solid adsorbents. Report No.2. Experimental verification of the equation of the isotherm of adsorption from solutions.

Izv. AN SSSR Otd.khim.nauk no.2:216-222 F 162.

(MIRA 15:2)

1. Institut fizicheskoy khimii AN SSSR.

(Adsorption)

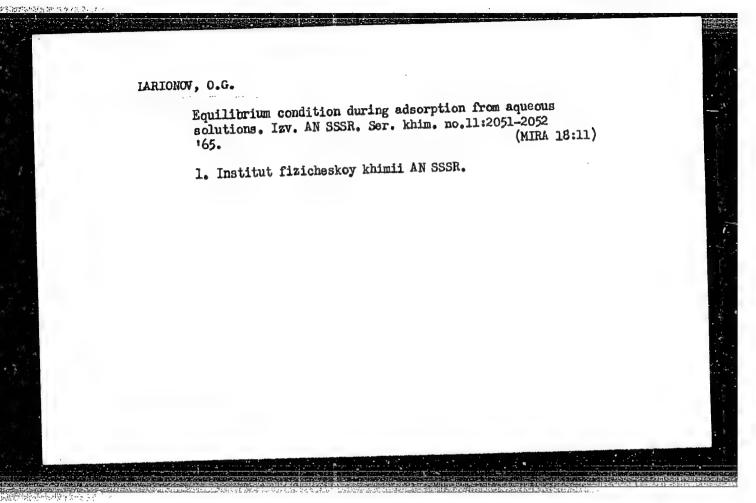


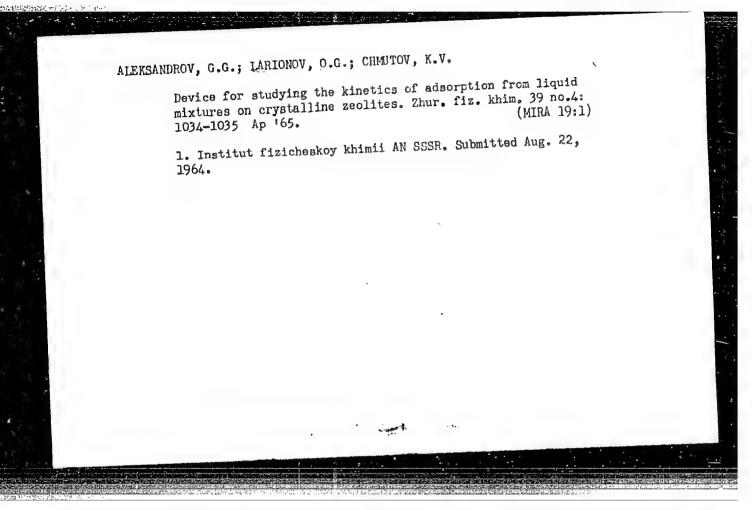
YELOVIN, S.Yu.; LARIONOV, O.G.

Application of mass-action law to adsorption equilibrium.

Izv.AN SSSR.Otd.khim.nauk no.3:531-533 Mr '62. (MIRA 15:3)

1. Institut fizicheskoy khimii AN SSSR.
 (Activity coefficients) (Phase rule and equilibrium)
 (Adsorption)





Calculating the true adsorption of mixture components from nonelectrolyte solutions. Zhur. fiz. khim. 39 no.9:2226— (MIRA 18:10) 2231 S. 165.

1. Institut fizioheskoy khimii AN SSSR.

CIA-RDP86-00513R000928710007-7 "APPROVED FOR RELEASE: 06/20/2000

 $\mathcal{O}_{\mathcal{N}}$ ONOV.

AUTHORS: Murin, A. N., Nefedov, V. D., Sinotova, Ye. N., 78-1-33/43

Larionov, O. V.

The Separation of the Nuclear Isomers of Tellurium, TITLE:

Mercury and Tin (Razdeleniye yadernykh izomerov tellura,

rtuti i olova)

PERIODICAL: Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, Nr 1,

pp. 181-183 (USSR)

ABSTRACT: After giving a review of the separation methods of the

> nuclear isomers of tellurium (references 1,2) and after their discussion the authors chose dimethyl-dinitrate of tellurium as the initial compound for the separation of the nuclear isomers of T127. It must be expected that the transition to an intermediate level will occur by means of an internal conversion and for this reason will be accompanied by a disturbance of the chemical binding of tellurium in the initial compound. Therefore a considerable portion of the nuclei of Te127 will be present as most simple anorganic

Card 1/4

forms in the ground state in the preparation dimethyl-dimitrate of tellurium. Te¹²⁷ in its ground state was isolated

The Separation of the Nuclear Isomers of Tellurium, Mercury and Tin

78-1-33/43

by means of the adsorption of these anorganic forms by ferric hydroxide. The extraction with isopropylether from 9 n HCl was intended for the removal of the an isotropic carriers, that is to say, iron. From the decay curve of the lowest isolated isomer (figure 1) follows, that only one tellurium isotope was existent, which had a half life of 9'3 hours. This testified to the presence of only the lowest isomer in the preparation. The yield of Te127 was determined to 80%, if it was accumulated in crystals, and to 94%, if it was accumulated in a solution. The latter value is in good correspondence with the known fact, that the isomeric transition in Te127 is converted to practically 100%. This implies, that the initial molecule is destroyed by every process of isomeric transition, which is accompanied by an internal conversion. The yield is somewhat lower, if accumulation takes place in crystals. The isolated radioactive Te127 predominantly takes its four-valent form and only 6 % of it take the six-valent one. This method possesses several advantages in comparison to the ones known hitherto (reference 1). If mercury is irradiated with neutrons according

Card 2/4

The Separation of the Nuclear Isomers of Tellurium, Mercury and Tin

78-1-33/43

to the reactions (n,7) and (n,2n), radioactive isotopes are formed: Hg197, Hg199, Hg205 and Hg205. Because at least six days elapsed until the separation was performed it can be assumed, that in the synthesized initial preparation -mercury diethyl only Hg²⁰³, Hg^{197m} and Hg¹⁹⁷ were present. From the investigations of the Laboratory for Radiochemistry of the University Leningrad (reference 3-6) it results, that the complete aliphatic mercury derivatives may undergo an irreversible destruction of the chemical bondings on isomeric transitions. The isolation of Hg197 in the ground level was performed by means of adsorption on manganese dioxide. The separation from the carrier can be achieved by methods, which are based on the volatility of mercury and its derivatives. The separation of the nuclear isomers as such can be determined from a comparison of the curves of decreasing activity of the mercury preparations (figure 2). When tin is irrariated by thermal neutrons, radioactive nuclei are formed: Sn113 (yields In113m, Sn117m and Sn119m by decay). From the three latter ones stable isotopes are produced by an isomeric transmutation: Sn117 and Sn119, Sn121, Sn123 and Sn125 were isolated in the ground

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The Separation of the Nuclear of Tellurium, Mercury and Tin

78-1-33/43

state from a benzene solution of stannic tetraphenyl by way of extraction. Because of the fact, that the isomers Sn123 and Sn125 have no genetic inter-relation, Sn121 and In113m will pass over into the water layer during the extraction. For this reason the activity measurement was started after the lapse of from 10-12 half life periods of In113m (T = 105 minutes). The decay curve of Sn121 is represented by figure 3. The accumulation of Sn121 with time was examined (figure 4) for the purpose of proving the genetic relation between Sn121 in ground state and Sn121m. The method described here may be considered the most universal. It makes furthermore possible to isolate the nuclei in a low isomeric state without carriers.

There are 4 figures, and 6 references, 4 of which are Slavic.

SUBMITTED:

June 18, 1957

AVAILABLE:

Library of Congress

Card 4/4

24 (5) AUTHORS:

Baranovskiy, V. I., Larionov, O. V.,

SOV/54-59-2-4/24

Nikitin, M. K., Tkachenko, A. A.

TITLE:

On the Problem of Natural Neutron Activity of Arsenic and Antimony (K voprosu o yestestvennoy neytronnoy aktivnosti

mysh'yaka i sur'my)

PERIODICAL:

Vestnik Leningradskogo universiteta. Seriya fiziki i khimii, 1959, Nr 2, pp 25-26 (USSR)

ABSTRACT:

In the papers by A. Dorabialska and M. Serwinski (Refs 1-3), it had been asserted that ordinary arsenic and antimony are sources of quick neutrons. By means of these neutrons, the authors had succeeded in activating Cu, Br, J and other elements. They set up a conversion scheme which, however, disagrees with the experimental mass determinations of the elements occurring in this scheme; even the inverse reactions had been observed in experiments. In order to prove that no neutrons are radiated from the said elements under natural conditions, the same experiments as described in the papers (Refs 1-3) were repeated in this paper. The exposition of the materials to be activated was carried out both by direct contact of As and Sb of high purity with activated materials,

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On the Problem of Natural Neutron Activity of Arsenic and Antimony

SOV/54-59-2-4/24

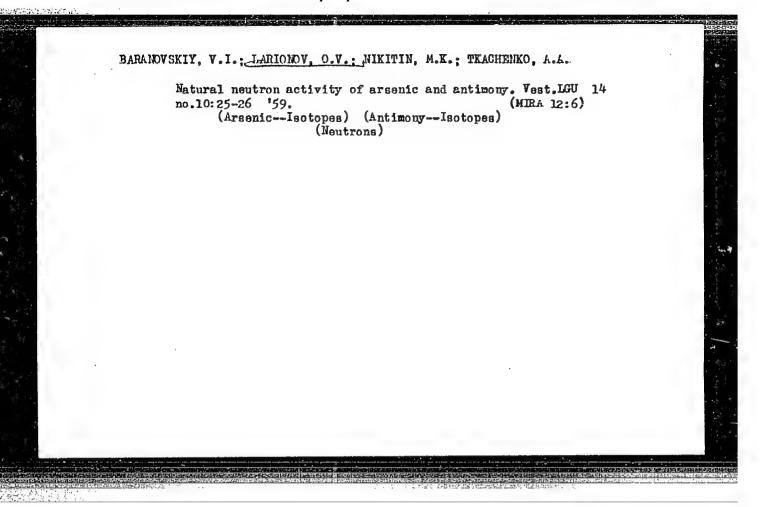
and with the use of moderators. All investigations proceeded with a negative result. Under experimental conditions as they were used in this investigation, a neutron decay of the As- and Sb-nuclei could have been detected only at a half-life period of $T_1 \neq 10^{16}$ a. For the self-activation of

the said nuclei, the background of the neutron capturing cross section should have been increased which has not been detected either. G-particles from a G-decay with energies > 0.05 Mev were missing. In all results obtained, the authors could not find a foundation for the assertion of a possible independent neutron decay in the As- and Sb-nuclei. Finally, the authors thank V. D. Nefedov for the discussions. There are 6 references, 1 of which is Soviet.

SUBMITTED:

June 14, 1958

Card 2/2



s/048/60/024/007/001/011 во19/во60

24,6600

AUTHORS:

Bashilov, A. A. (Deceased), Larionov, O. V., Nikitin,

Smirnov, V. B. M. K.,

TITLE:

Eu 145 Production in Ta Spallation Reactions

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, 1960,

Vol. 24, No. 7, pp. 788-790

TEXT: This, is the reproduction of a lecture delivered at the 10th All-Union Conference on Nuclear Spectroscopy held in Moscow from January 19 to 27, 1960. The authors studied the Eu145 production in Ta spallation reactions produced by 660-Mev protons. The synchrocyclotron used belonged to the OIYaI (Joint Institute of Nuclear Research). The Eu isotopes produced in the reactions were examined with gamma rays. Six hours after the Ta target irradiation, the rare earths were chemically separated and the fractions of the individual rare earth elements were further separated. The Eu fraction was purified chromatographically and was then added to a diluted HNO3 solution containing La2+ for the prevention of absorption.

Card 1/2

Eu 145 Production in Ta Spallation Reactions

S/048/60/024/007/001/011 B019/B060

Gamma emission was investigated with a NaI scintillation spectrometer. Apart from the gamma lines of Eu¹⁴⁶, Eu¹⁴⁷, and Eu¹⁴⁹, the authors identified 0.89, 1.66, 1.86, and 2.0 Mev lines, whose intensity drop corresponded to a half-life T = 5.5 days (Fig. 2). Thorough examinations made on the daughter products gave evidence that the observed Eu activity with a half-life of 5.5 days must be ascribed to the Eu¹⁴⁵ isotope. The data obtained here agree with those of Hoff and others (Ref. 4). The authors thank I. B. Stankevich for having conducted the chemical operations, and V. B. Savichev for his assistance in the measurements. There are 4 figures and 4 references: 3 Soviet and 1 US.

ASSOCIATION:

Nauchno-issledovatel'skiy fizicheskiy institut Leningradskogo gos. universiteta im. A. A. Zhdanova (<u>Scientific Research Institute of Physics of the</u> Leningrad State University imeni A. A. Zhdanov)

Card 2/2

S/048/60/024/007/018/032/XX B019/B056

24,6720 AUTHORS:

الماسي سرو

Grigor'yev, Ye. P., Larionov, O. V., Nikitin, M. K.,

Sakharov, S. L., and Sergeyev, V. O.

The Determination of the Halflife of \underline{Dy}^{159} , \underline{Ho}^{160} , \underline{Tu}^{166} TITLE:

and Lu 173

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, 1960, PERIODICAL:

Vol. 24, No. 7, pp. 841-844

TEXT: This paper was read at the 10th All-Union Conference on Nuclear Spectroscopy, which took place from January 19 to January 27, 1960 at Moscow. The isotopes investigated were obtained by the irradiation of Tatargets with 660-Mev protons in the synchrocyclotron of the Ob"yedinennyy ir stitut yadernykh issledovaniy (Joint Institute of Nuclear Research) and a subsequent chemical and chromatographical separation. For determining the halflife an end-window counter was used, which was protected by a Pb-shield.

As a control isotope, Dy 159 was selected. The authors determined a halflife $T = 139 \pm 10$ days, which agrees with the data obtained by other

Card 1/2

24.6720 AUTHORS:

s/048/60/024/007/019/032/XX B019/B056

Grigor'yev, Ye. P., Larionov, O. V., Nikitin, M. K., Sakharov, S. L., and Sargeyev,

TITLE:

The Y-Spectra of the Isotopes of the Tantalum Fraction

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, 1960, Vol. 24, No. 7, pp. 845-846

This paper was read at the 10th All-Union Conference on Nuclear Spectroscopy, which took place from January 19 to January 27, 1960 at Moscow. In the synchrocyclotron of the Olyal, a Ta-target was irradiated with 660-Mev protons, following which, tantalum was separated and the radioactive Ta-isotopes were investigated by means of an automatic scintillation-8-spectrometer. According to the halflife of the &-lines, the Ta-isotopes may be subdivided into two groups. There are some isotopes with a halflife T of roughly 8 hours, and others with T = 53 hours. The energy and the relative intensities of the y-lines of those Tamisotopes whose T is about 8 - 11 hours, are given in Table 1; Card 1/2

The y-Spectra of the Isotopes of the Tantalum S/048/60/024/007/019/032/XX Fraction S/048/60/024/007/019/032/XX

 E [kev]
 55
 115+5
 210+10
 270
 350
 500
 1150
 1700

 I_f
 100
 10
 7
 2
 2
 0.3
 0.7
 0.3

By comparison with data obtained by other authors, the authors draw the conclusion that in their Ta-fraction the isotopes Ta¹⁷⁶ (8 hours) and

Ta¹⁷⁵ (11 hours) are present. In Table 2, the energies and the relative intensities of the hard y-lines of the Ta-isotope of a halflife of 8 hours are given:

 Er
 [Mev]
 1.7
 2.2
 2.3
 2.7
 2.8

 I₂
 3
 1
 0.3
 1

These hard lines may possibly belong to a Ta 176 -decay. From the data obtained here, the authors conclude that the mass difference between

Ta 176 and Hf 176 is more than 3 Mev. There are 2 figures, 2 tables, and 7 references: 1 Soviet and 6 US.

ASSOCIATION: Nauchno-issledovatel'skiy fizicheskiy institut Leningradskogo gos. universiteta im. A. A. Zhdanova (Scientific Research

Institute of Physics of Leningrad State University imeni

A. A. Zhdanov)

Card 2/2

21:066

s/054/61/000/002/004/005 B101/B207

5.2300

1087

Larionov, O. V., Nikitin, M. K.

AUTHOR: TITLE:

The problem of separating rare-earths elements from

tantalum

PERIODICAL:

Leningradskiy Universitet. Vestnik. Seriya fiziki i

khimii, no. 2, 1961, 73 - 76

TEXT: The aim of the present study was the chromatographic partition of rare-earths elements (REE) from tantalum without using a carrier, in order to obtain tantalum completely free from REE isotopes. The usual precipitation of REE as fluorides is incomplete. Furthermore, other elements, such as Hf, are coprecipitated. Thus, the separation and study of the radioactive isotopes of Hf are rendered difficult. On the assumption that REE exist in a Hf medium as cations, tantalum (as well as Hf and Zr) as anions (TaF_7^2 , TaF_8^3) the adsorption of REE cations on cation

exchangers was studied. The distribution of microquantities of Eu Card 1/4

21,066

S/054/61/000/002/004/005 B101/B207

The problem of separating ...

among the Hf solution, the resins KY-2 (KU-2), and Dowex-50 was investigated: Eu¹⁵² was completely adsorbed at concentrations of 10⁻⁷ - 10⁻⁸ g/ml of 5-7 mg resin, while Ta¹⁸² was not adsorbed. Further experiments were carried out at room temperature with a chromatographic column of 2 mm diameter, filled with commercial KU-2 in the form of H (grain size, 50 µ; layer height, 3-4 mm), on plexiglass wadding. The solution was pressed through the exchanger by a mercury column. Eu¹⁵² was completely separated from the inactive Ta (concentration up to 0.35 g/ml) at a rate of one drop every 10-12 sec (1 drop ~ 1/20 ml). At a higher flow rate (one drop every 2-4 sec), up to 10% of Eu was not adsorbed by the exchanger. The presence of HNO₃ deteriorated the results. The elution curves of Fig. 2 show that concentrated HNO₃+saturated H₃BO₃ solution is the most effective eluent. The method described was successfully applied to the quantitative separation of REE isotopes from tantalum which was irradiated with Card 2/4

21,066

The problem of separating ...

Card 2/4

S/054/61/000/002/004/005 B101/B207

among the Hf solution, the resins KY-2 (KU-2), and Dowex-50 was investigated: Eu¹⁵² was completely adsorbed at concentrations of 10⁻⁷ - 10⁻⁸ g/ml of 5-7 mg resin, while Ta¹⁸² was not adsorbed. Further experiments were carried out at room temperature with a chromatographic column of 2 mm diameter, filled with commercial KU-2 in the form of H (grain size, ~50 \mu; layer height, 3-4 mm), on plexiglass wadding. The solution was pressed through the exchanger by a mercury column. Eu¹⁵² was completely separated from the inactive Ta (concentration up to 0.35 g/ml) at a rate of one drop every 10-12 sec (1 drop ~ 1/20 ml). At a higher flow rate (one drop every 2-4 sec), up to 10% of Eu was not adsorbed by the exchanger. The presence of HNO₃ deteriorated the results. The elution curves of Fig. 2 show that concentrated HNO₃+saturated H₃BO₃ solution is the most effective eluent. The method described was successfully applied to the quantitative separation of REE isotopes from tantalum which was irradiated with

21,066 s/054/61/000/002/004/005 B101/B207

The problem of separating ...

660-Mev protons. This mode of separation is more complete than that performed by the fluoride method. There are 2 figures and 4 references: 1 Soviet-bloc and 3 non-Soviet-bloc. The 3 references to English-language publications read as follows: H. J. Hettel, V. Fassel, Anal. Chem., 27, 1311, 1955; Nervik a. Seaborg, Phys. Rev., 97, 1092, 1954; H. Gest, W. H. Burgus, T. H. Davies, Radiochemical studies. The fission products. Book 1, paper 13, 1951

SUBMITTED: May, 1959

Fig. 2: Curves of REE elution. Legend: 1) HNO3; 2) HCl; 3) HNO3+ H3BO3; a) pulses per min., b) number of drops.

Card 3/4

CIA-RDP86-00513R000928710007-7" APPROVED FOR RELEASE: 06/20/2000

S/186/61/003/001/016/020 A051/A129

21.3200 AUTHORS: Murin, A.N., Nefedov, V.D., Larionov, O.V.

TITLE: The separation of nuclear isomers of tellurium

PERIODICAL: Radiokhimiya, v 3, no 1, 1961, 90-96

TEXT: The authors have developed a new method for the separation of nuclear isomers of tellurium and the separation of lower isomer compound states without a carrier, as well as a method for the separation of radio-chemically pure Te127 from irradiated tellurium dimethyldinitrate with neutrons (and pure Te127 from irradiated tellurium of Te127 from the irradiated quanta). They show that the extraction of Te127 from the irradiated sample reaches a yield close to 100%, which corresponds to the break of the chemical bond in each converted isomer transition. The greater part (about chemical bond in each converted isomer transition. The greater part (about 91%) of the extracted Te127 is in the lower tetra-valent state and only 91%) of the extracted Te127 is in the lower tetra-valent state and only separation of the main isomer state of tellurium was tellurium dimethyldiseparation of the main isomer state of tellurium was tellurium dimethyldiseparation.

Card 1/6

The separation of nuclear isomers of tellurium

S/186/61/003/001/016/020 A051/A129

nitrate (CH₃)₂Te(NO₃)₂. The latter was formed from tellurium dimethyldiciodine: Te + 2(CH₃)I (CH₃)₂TeI₂. The authors investigated various ways of isolating Te in the basic state: 1) extraction of the basic salts of Mn on the residue formed when an alcohol solution of Mn(CH₂COO)₂ is added to the acetone solution (CH₃)₂Te(NO₂)₂ was found to be inconvenient, since the residue retained most of the initial quantity of the compound; 2) extraction on the residue of H₂WO₄ gave a small yield; 3) extraction on MnO₂ was impossible due to oxidation of the initial compound and dissolution of MnO₂; in the basic state. The authors adopted the Te isolation method on iron hydroxide. The curve of Fig 3 shows that there is only one isotope with a half-life of 9.3 hours, which proves the presence of Te¹²⁷ in the sample in the basic state. The degree of impurities was studied using metastable Turther, the radiochemical purity of samples was investigated (Fig 2). Further, the radiochemical purity of samples produced according to the authors methods was compared to that produced according to the methods of Siborg, Livinhood and Kennedy. The average yield was found to be 79.5±2.2%

Card 2/6

The separation of nuclear isomers of tellurium

S/186/61/003/001/016/020 A051/A129

when accumulated in crystals. The high yields noted by the authors are thought to be the result of the sharply expressed irreversibility of the occurring chemical changes during isomer transition when using (CH,), Te(NO,) The data of Table 2 show that with an accumulation of Te in the crystals the yield of the basic state is somewhat less since in this case there is a greater stability of the basic state of Te 127 in the form of the initial tellurium dimethyldinitrate compound. The difference in the chemical behavior of the tetra and hexa-valent states of Te helps to solve the problem of Te distribution between these valency states. The study of this question was carried out by the isotopes carrier method corresponding to various chemical compounds (TeO2 and H2TeO4). The separation of the 6- and 4-valent Te was based on the reduction of the latter to the elemental state by sulfurdioxide in a 3 n solution of HCl (Ref 12). The average yields are 127 equal to 8.511.2% and 91.511.2%, respectively. The fact that most of Te is in the lower valency state is explained by secondary processes which occur after the above-mentioned phenomena. The activation of Te in the main state was conducted on a betatron and the separation of Te in the main state was carried out according to the reaction (γ,n) (Fig 4). There are 4 figures, 4 tables and 14 references: 6 Soviet-bloc, 8 non-Soviet-bloc. Card 3/6

S/186/61/003/005/020/022 E111/E485

AUTHORS: Nefedov, V.D., Larionov, O.V.

TITLE: A constant carrierless T1206 source

PERIODICAL: Radiokhimiya, v.3, no.5, 1961, 639

TEXT: The authors describe a method of production of preparations of $T1^{206}$ without carrier. This method can also be used for detecting the presence of Bi^{210} in bismuth preparations. It is known that if Bi^{210} obtained by neutron irradiation of bismuth is introduced into an organometallic compound $\left[(C_6H_5)_3Bi\right]$ or $(C_6H_5)_7PiCl_2$, this preparation can serve as a carrierless $T1^{206}$ source. The metallic bismuth was neutron irradiated for a long time and then kept for half a year. After this it was dissolved and repeatedly cleaned from polonium. As most convenient starting compound, $(C_6H_5)_3BiCl_2$ was chosen being more stable than $(C_6H_5)_3Bi$. In this case, $T1^{206}$ was extracted from the benzene solution $(C_6H_5)_3Bi^*Cl_2$ by 5% hydrochloric acid. In the case of $(C_6H_5)_3Bi^*T1^{206}$ was extracted from the ethereal solution of the bismuth organic compound by water. It was found that the yield of $T1^{206}$ was higher if it was directly separated on MnO_2 from the benzene solution by addition of a few drops of Card 1/2

A constant carrierless Tl²⁰⁶ source

S/186/61/003/005/020/022 E111/E485

 ${\rm KMnO_4}$ in acctone and hydrogen peroxide. This proved that part of the resulting ${\rm Tl^{206}}$ is in the form of organo-metallic compounds. To identify ${\rm Tl^{206}}$ from the half-life period it was isolated from the enriched fraction with an isotope carrier in the form of ${\rm Tl(OH)_3}$ or by adsorption on ${\rm MnO_2}$. The half-life period T was found from the equation

 $T = -t \frac{0.301}{\log A_{2t} - \log A_{t}}$

where t is a definite time interval at which activity determinations were made. At is the number of disintegrations in time t and A2t that in time 2t. T values (minutes) obtained were 4.5, 4.8, 4.9. There are 1 figure and 5 references: 2 Soviet-bloc and 3 non-Soviet-bloc. The three references to English language publications read as follows: Ref.1: H.M.Neumann, J.J.Howland, I.Perlman, Phys. Rev., 77, 720 (1950); Ref.2: H.B.Levy, I.Perlman, Phys. Rev., 85, 758 (1952); Ref.3: H.B.Levy, I.Perlman, Phys. Rev., 94, 152 (1954).

Card 2/2

S/048/61/025/001/016/031 B029/B060

24.6720

Berlovich, E. Ye., Larionov, O. V., Tunimanova, E. N.,

Khay, D. M.

TITLE:

Study of the decay schemes of Gd 146, Gd 147, and Gd 149 by a

beta - gamma coincidence spectrometer

PERIODICAL:

Izvestiya Akademii nauk SSSR. Seriya fizicheskaya, v. 25,

no. 1, 1961, 90-97

TEXT: A study has been made of the cascade properties of transitions in gadolinium isotopes by the method of coincidences with a view to defining the details of the decay schemes of these isotopes. N. M. Anton'yeva, A. A. Bashilov et al. (Refs. 2,3,4), in their papers submitted to the 8th All-Union Conference on Nuclear Spectroscopy of 1958, had offered a thorough study of the spectra of conversion electrons of Gd¹⁴⁶, Gd¹⁴⁷, and Gd¹⁴⁹. B. S. Dzhelepov, V. A. Sergiyenko et al. (Refs. 5,6) studied the coincidences between the conversion electrons of these isotopes in 1959. Fig. 2 shows the block diagram of the coincidence spectrometer,

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89250 S/048/61/025/001/016/031 B029/B060

1×

Study of the decay schemes of ...

consisting of two branches, used here. The two branches represent a sector-type magnet spectrometer with improved focusing and a scintillation spectrometer with a NaI crystal. The recorders were two time photomultipliers of the type \$\phi 3\forall -14 (FEU-14) after G. S. Vil'dgrube. Measurement results: Gd 146: Fig. 3 shows the curve of the coincidences of electrons of the K line of transition (114.8 + 115.5)kev with the gamma rays of the gadolinium fraction. The measurements took place 100 days after the separation of the fraction from the target irradiated with 660-Mev protons. Fig. 4 shows the analogous curve for the K line of the 155-kev transition. Two incompletely resolved coincidence peaks are observed; peak 1 characterizes the coincidences K114.8 - y115.5 and 115.5 - γ 114.8; peak 2 refers to K(114.8 + 115.5) - γ 155. The results found, while confirming the cascade property of all of the three transitions, do not, however, add any new information to the results given by B. S. Dzhelepov and V. A. Sergiyenko (Ref. 5). Still, they may be regarded as a good confirmation of the hitherto assumed decay scheme of Gd 146. Fig. 5 shows the peaks of the coincidences of the 229-kev electrons with the gamma rays (scintillation branch), and Fig. 6 shows the

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89250 S/048/61/025/001/016/031 B029/B060

Study of the decay schemes of ...

peaks of the coincidences of the 396 kev-K-line electrons with the same gamma rays. The K229 electrons coincide with the 396-kev gamma rays. The peak of coincidences is, however, widened by an admixture of 370-kev gamma quanta. In the spectrum of coincidences there are still further, although not sharp, maxima, which correspond to the 560 ±20, and 760±25-kev energies, as well as a poorly marked coincidence peak in the 900-kev range. Weak maxima are also observed with 396-kev electrons, namely, in the 480 ± 30 and 560 ± 30 -kev energy range. Figs. 7 and 8 show the coincidence curves of conversion K electrons of the 149.8 and 346-kev transitions with the gamma rays recorded in the scintillation branch. K 149.8 electrons provide coincidences with the 346 and 530 20-kev gamma quanta. K 346 electrons provide coincidences with 150 and 298-kev gamma quanta. According to the results obtained, the 298-kev transition in the nucleus of 63Eu 149 is surely to be found in the 346- and 149.8-kev gamma cascade. This transition lies above the isomeric level and proceeds from the 795-kev level. Spin and parity 9/2 or 11/2 must be ascribed to this level. The intensities of 346 and 298-kev transitions are almost equally high. The excitation of the 497-kev level by electron capture is,

Card 3/9

Study of the decay schemes of ...

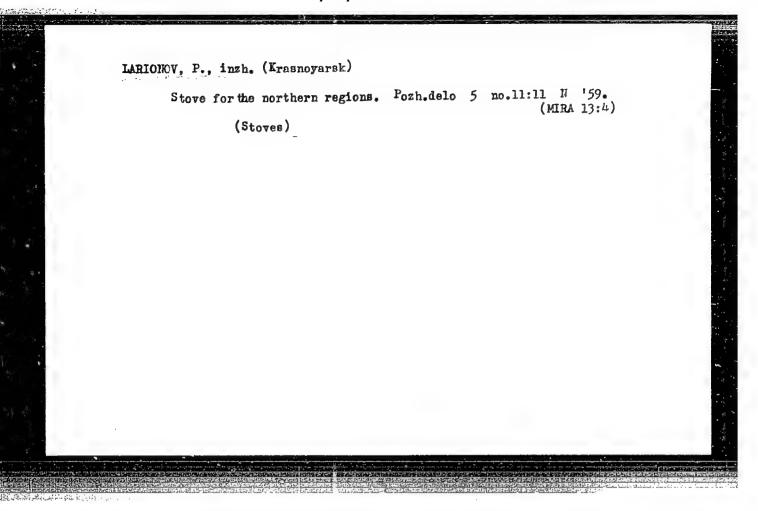
S/048/61/025/001/016/031 B029/B060

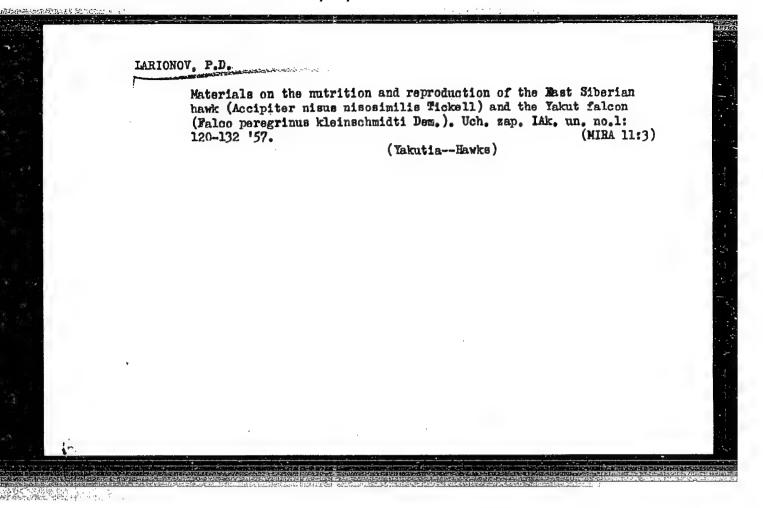
in fact, more probable than the excitation of the 795-kev level. The other results found regarding the coincidences are in good agreement with the decay scheme of Gd 49 suggested by N. M. Anton'yeva et al. (Ref. 3). The article under consideration is the reproduction of a lecture delivered at the 10th All-Union Conference on Nuclear Spectroscopy, which took place in Moscow from January 19 to 27, 1960. There are 11 figures, 1 table, and 9 references: 8 Soviet-bloc and 1 non-Soviet-bloc.

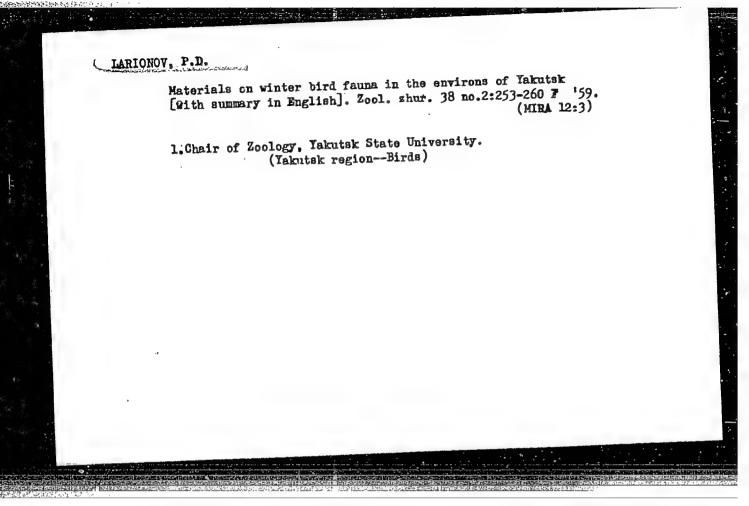
ASSOCIATION: Fiziko-tekhnicheskiy institut im. A. F. Ioffe Akademii nauk SSSR (Institute of Physics and Technology imeni A. F. Ioffe, Academy of Sciences USSR)

Legend to Fig. 2: 1) magnetic spectrometer (a) source container, (6) deflection chamber, (6) counter chamber, (2) source; 2) limiters; 3) variable delay line; 4) fast-coincidence block, (4) amplifier; 5) differential pulse height analyzer; 6) triple coincidence circuit; 7) counter.

Card 4/9







LARIONOV, P.D.

Conditions of hibernation of vipers at the northern limit of their range in the Lena Valley. Zool. zhur. 40 no. 2:289-290 F '61. (MIRA 14:2)

1. Department of Zoology, State University of Yakutsk. (Kytyl-Zhura region-Serpents) (Hibernation)

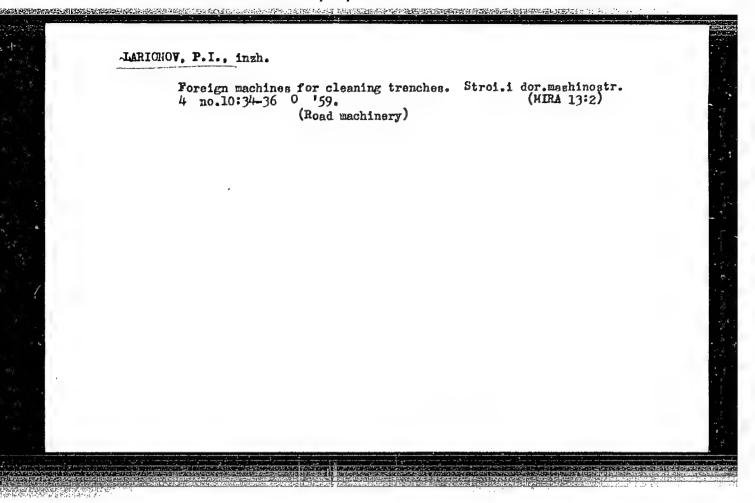
MIKHAYLOV, N.V.; BUKOV, G.A.; GORBACHEVA, V.O.; MAKAROVA, T.P.; v rabote prinimali uchastiye: Larionov, P.E.; Sorokina, V.I.; Zotov, Ya.E.

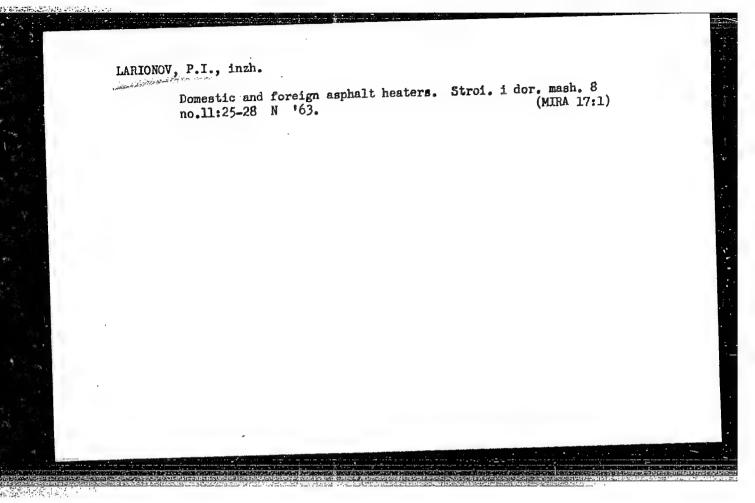
Studying the formation mechanism of synthetic fibers from molten materials. Khim.volok. no.1:33-36 *59. (MIRA 12:8)

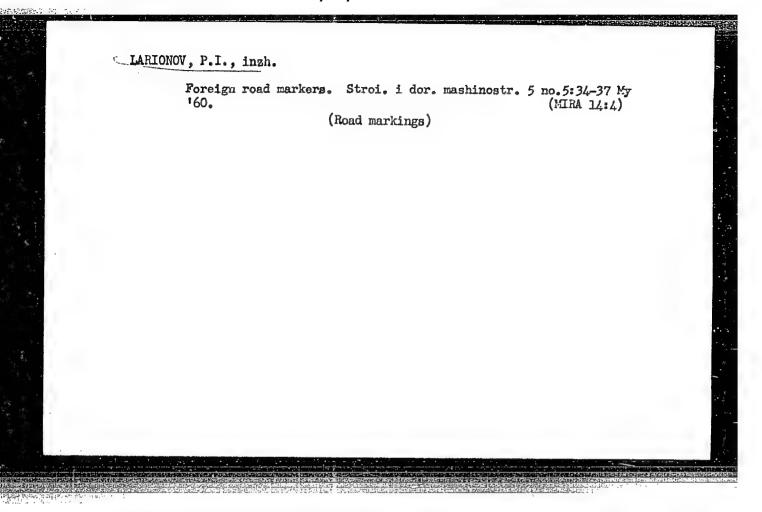
1. Vsesoyuznyy nauchno-issledovatel'skiy institut iskusstvennogo volokna.

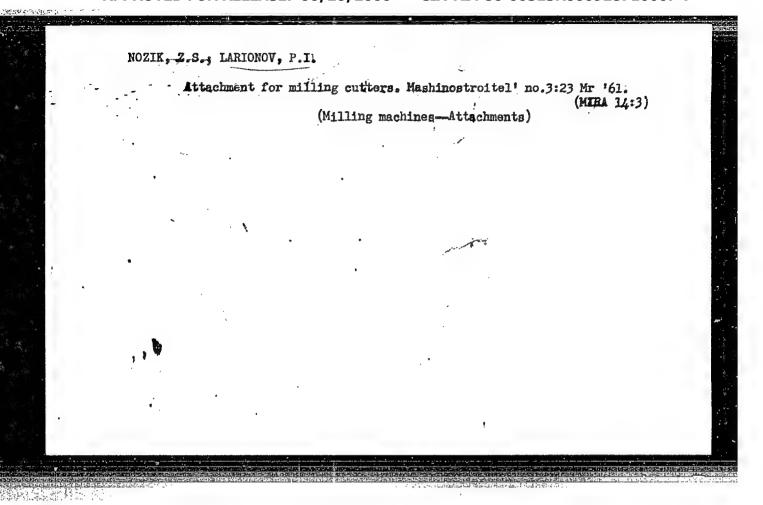
(Textile fibers, Synthetic)

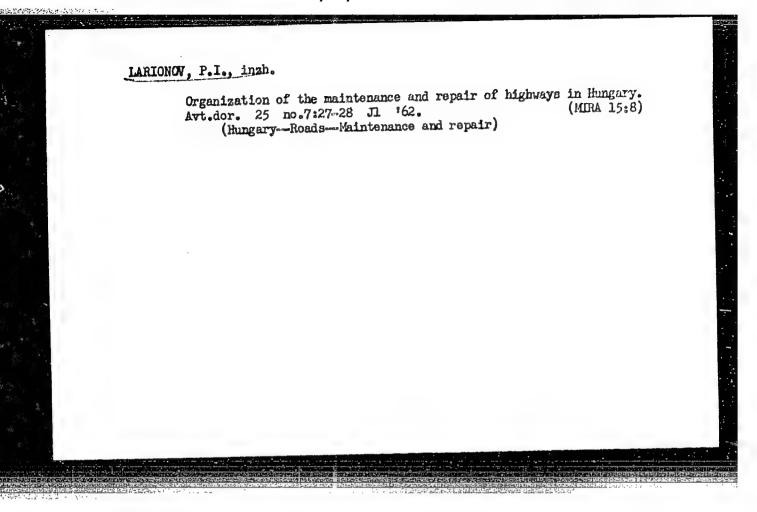












MIKHAYLOV, N.V.; SHEYN, T.I.; GORBACHEVA, V.O.; TOPCHIBASHEVA, V.N.;

v rabote prinicali uchastiye tekhniki-laboranty; IARIONOV, P.M.;

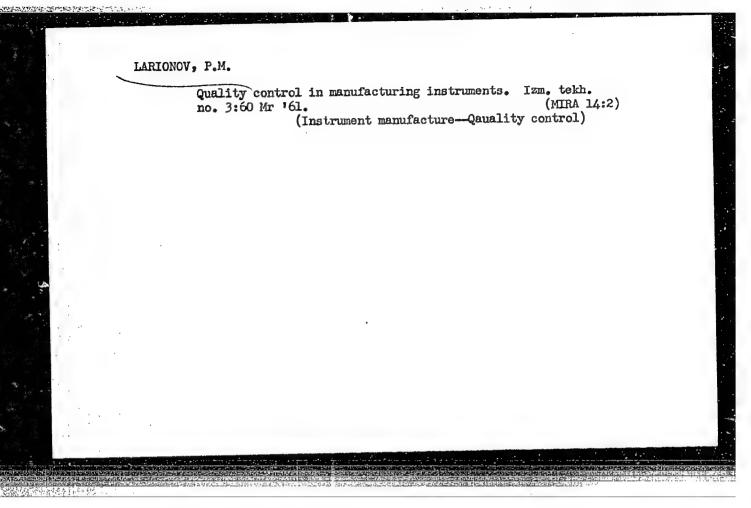
VLASOVA, L.P.; MURASHKINA, S.I.

Investigating the molecular structure of synthetic fibers.

Part 14: Physicochemical and physicomechanical properties of the polycapramide - polyundecanamide polyamide group. Vysokomsoed. 1 no.2:185-190 F *59. (MIRA 12:10)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut iskusstvennogo volokna.

(Textile fibers, Synthetic) (Amides)



SHCHEGLOVA, O. P., kand. fiz.-matem. nauk; LUT, B. F.; MECHITOV, I. I., kand. tekhn. nauk (Tbilisi); IVERONOVA, I. M., kand. geograf. nauk (Moskva); IOGANSON, V. Ye. (Moskva); LARIONOV, P. M. (Uzhgorod)

Mud torrents, Prioroda 52 no.1:90-96 63. (MIRA 16:1)

1. Tashkentskiy gosudarstvennyy universitet im. V. I. Lenins for Shcheglova). 2. Baykal'skaya limnologicheskaya stantsiya, poselok Listvenichnoye, Irkutskaya obl. (for Lut).

(Runoff) (Erosion)

IARIONOV, S.F., inzh.; POZDNYAK, A.A., inzh.

Concerning I.I.Khazovskii's article "Change-over to centralized structure of the departments in electric power plants." Elek. sta. 33 no.7:90-91 J1 '62. (MIRA 15:8) (Electric power plants) (Khazovskii, I.I.)

LARIONOU, S.N.

USBR/General and Specialized Zoology. Ensects. Injurious Insects and Ticks. Comeral Problems

Abs Jour : Ref Zhur - Biol., No 11, 1958, No 49536

: Larionov S.H. Author

: Lethrus Bootles as Pests of Agricultural Croys in Inst

Southern Kazakhstan. Titlo

Orig Pur: Zashchita rast. ot vredit. i boleznoy, 1957,

No 4, 52

Abstract: Lethrus bituberculatus and L. scoparius were found in quantities of 0.1-8 individuals per lm2 in regions of the former Golodnaya Steppe on virgin lands, waste lands, soddy shoulders of roads and irrigators, old uncultivated alfalfa

fields and vineyards. They damaged tomatoes, cabbage, carrots, corn, cotton and grape vine. Methods of control were tilling and stirring up

: 1/2 Card

12

Larionov, S.P.

137-1957-12-23701

Translation from: Referativnyy zhurnal, Metallurgiya, 1957, Nr 12, p 123 (USSR)

AUTHOR:

Larionov, S.P.

TITLE:

An Investigation of the Process of Upsetting Cylindrical Specimens With an Opening (Issledovaniye protsessa osazhivaniya tsilindricheskikh obraztsov s otverstiyem)

PERIODICAL: Sb. stud. n auchn. rabot. Belorussk. politekhn. in-t, 1957, Nr 3, pp 5-8

ABSTRACT: The possibility of welding-up defects in metals was studied on specimens with openings (O). The changes in the shape of the O were investigated under monoaxial compression conducive to the complete disappearance of O. Complete welding-up would not occur, since almost always minute cracks remained at the point of the weld. Therefore, bars of non-plastic alloys of insufficiently dense structure should be extruded from the container through a die in order to transform the cast structure into a deformation structure under conditions most favorable for the climination of density discontinuities.

Card 1/1

1. Metals-Defects-Salvage methods

V. O.

S/078/61/006/001/005/019 B017/B054

AUTHORS: Batyayev, I. M., Larionov, S. V., Shul'man, V. M.

TITLE: Stability of Complex Compounds of Lanthanum, Cerium, Praseodymium, and Neodymium With Aspartic Acid

PERIODICAL: Zhurnal neorganicheskoy khimii, 1961, Vol. 6, No. 1, pp. 153 - 156

TEXT: Complex compounds of lanthanum, cerium, praseodymium, and neodymium were more precisely defined by potentiometric titrations of 0.01 molar solutions of aspartic acid with 0.1 N KOH in the presence and absence of rare earth ions. The pH value was measured at 25 °C by an JM-5 (LP-5) potentiometer and a glass electrode. Two series of titrations were conducted with a component ratio of $c_{\rm H_2A}$: $c_{\rm M}$ 3+ = 1: 1 and 2: 1

(H₂A = aspartic acid). The potentiometric titration curves are shown in Figs. 1 and 2. The stability of complex compounds of lanthanum, cerium, praseodymium, and neodymium with aspartic acid increases in the following Card 1/2

Stability of Complex Compounds of Lanthanum, S/078/61/006/001/005/019 Cerium, Praseodymium, and Neodymium With B017/B054

order: La < Ce < Pr < Nd. The authors conclude from their results that the interaction between La 3+, Ce 3+, Pr 3+, Nd 3+ and aspartic acid is not restricted to the formation of MA+ and MA2 complexes. There are 3 figures, 1 table, and 9 references: 2 Soviet, 2 US, 2 British, 2 Danish, and 1 Swiss.

SUBMITTED: September 2, 1959

Card 2/2

S/200/62/000/012/003/005 D204/D307

AUTHORS:

Batyayev, I.M. and Larionov, S.V.

TITLE:

Stability of the complexes of lanthanum, prascodymium and neodymium with glycocol

PERIODICAL:

Akademiya nauk SSSR. Sibirskoye otdeleniye. Izvestiya, no. 12, 1962, 69-73

The present paper is concerned with the study of the stability of complexes of La, Pr, and Nd with glycocol, and of Nd with a alamine and serine, since such work may be of importance in the study of the complexing of lanthanons with polypeptides. The stability constants were determined by potentiometric titrations, at 25 ± 0.05°C, of the above amino acids with 0.1 N KOH, in the presence and absence of the ions of La, Pr. Nd. The acid:metal ratio was 1:1 or 3:1. For a titration, 100 ml of 0.001 M amino acid were placed in the cell, followed by 0.3 to 1.7 ml of neutral MCl₂ (M = lanthanon) to an ionic strength of 0.1. The latter value did not rise by more than 5% during experiment. Logarithms of the 1st stab-

Stability of the complexes ...

S/200/62/000/012/003/005 D204/D307

ility constants (x1) were found to be respectively 4.18, 4.66 and 4.74 for La. Pr and Nd complexes with glycocol. A value of 4.3 was also found for log x2, for the glycocol complex of Nd. For complexes of Nd with α-alamine and serine, log x1 was respective-1y 5.04 and 4.52. Comparisom of these results with earlier work (Zh. neorg. khim., VI, 153 (1961); Izv. Sib. otd. AN SSSR, no. 2, and Nd with aspartic acid is higher than the stability of the complexes of La, Pr plexes of the same elements with glutamic acid or glycocol. There

ASSOCIATION:

Institut neorganicheskoy khimii Sibirskogo otdeleniya AN SSSR, Novosibirsk (Institute of Inorganic Chemistry of the Siberian Branch of AS USSR, Novosibirsk)

SUBMITTED:

January 20, 1962

Card 2/2

BATYAYEV, I.M.; LARIONOV, S.V.

Stability of complex compounds of lanthanum, cerium, praseodymium and neodymium with glutamic acid. Izv. Sib. otd. AN SSSR no.2:113-115 '62. (MIRA 16:10)

1. Institut neorganicheskoy khimii Sibirskogo otdeleniya AN SSSR, Novosibirsk.

IARIONOV, S.V.; SHUL'MAN, V.M.; PODOL'SKAYA, L.A.

Complex formation of nickel with o-thiosalicylic acid. Zhur.
neorg. khim. 9 no.10:2333-2338 0 '64.

1. Institut neorganicheskoy khimii Sibirskogo otdeleniya Akademii nauk SSR.

SHUL'MAN, V.M.; LARIONOV, S.V.; KRAMAREVA, T.V.; YEFREMOVA, T.D.

Oxido-reduction rotentials of the system thiourea - formamidine dissulfide in some mixed solvents. Izv. AN SSSR. Ser. khim. no.7:1257.

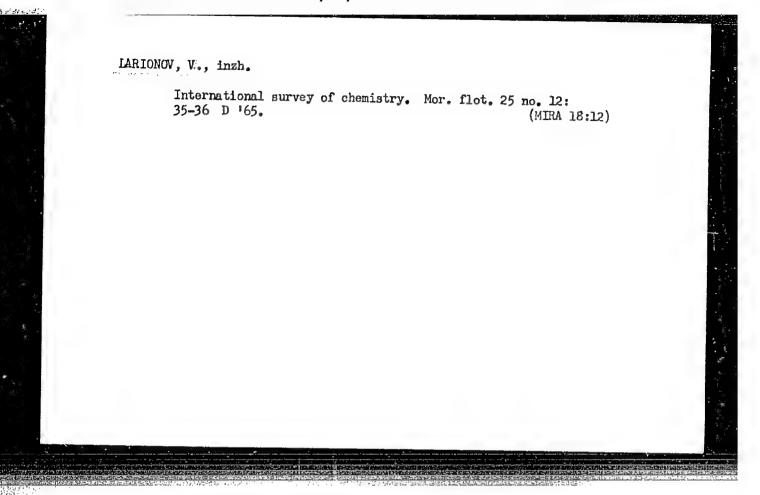
(MIRA 18:7)

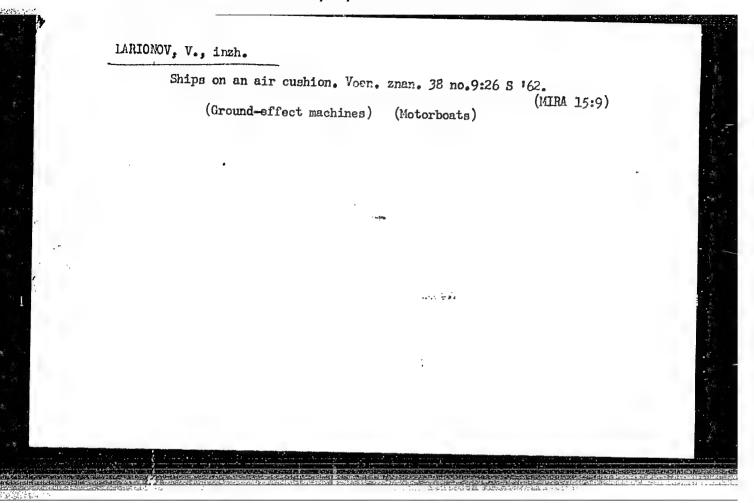
1. Institut neorganicheskoy khimit Sibirskogo otdeleniya AN SSSR.

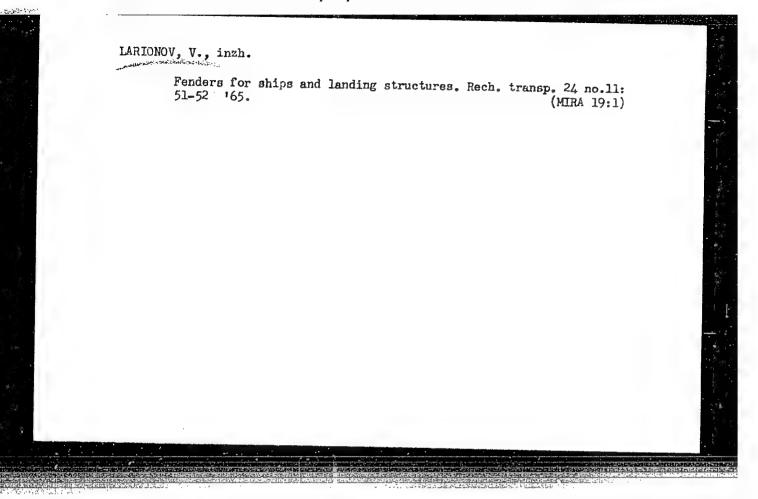
New marine evaporator systems. Mor. flot 22 no.10:43-44 0 '62.

(MIRA 15:10)

(Feed water purification—Equipment and supplies)







L 16905-65 EWT(m)/EWA(d)/EWP(t)/EWP(b) IJP(c)/ASD(f)-2/SSD/ASD(m)-3/AFTC(p)
ACCESSION NR: AP\$049180 MJW/JD/WB S/0314/64/000/005/0028/0029

AUTHOR: Pul'tsin, N. M. (Candidate of technical sciences); Larionov,
V. A. (Engineer)

Till: Investigation of titanium-alloy strength in an aggressive medium

SUJRCE: Khimicheskoya i neftyanoye mashinostroyeniye, no. 5, 1964,
28-29

TOPIC TAGS: fitanium, titanium alloy, alloy property, VT14 alloy,
AT3-ailcy, corrosion, stress corrosion, sulfuric acid

ABSTRACT: Tests have been conducted to determine the strength of the

ABSTRACT: Tests have been conducted to determine the strength of the VT14 and AT8 titanium alloys in an aggressive medium and to investigate the effect of the surface layer formed as a result of gas absorption during annealing at 880C for 0.5 to 2 hr. Alloys were stressed to 92.5, 95, or 97.5% of their tensile strenght, in 20% solfuring acid. AT8 alloy was found to be more resistant to the combined effect of stress and corrosion. Under a stress of 95% of the considerable strength, VT14 alloy failed in 30 min and AT8 alloy failed

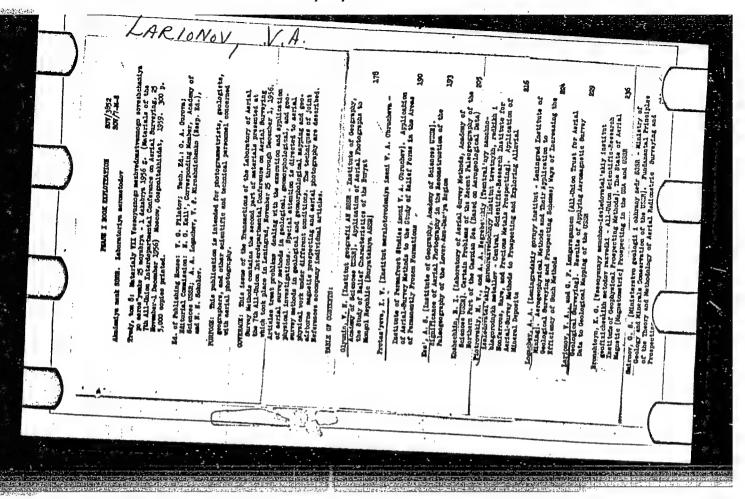
Card 1/2

L 16905-65 ACCESSION NR: AP4049180 in 3 hr 15 min. Under a stress of 92.5% of the tensile strength, neither alloy failed after 24 hr. The surface layer formed under the effect of heat treatment increases the strength of both alloys in the aggressive medium. In tests under a stress of 95% of the tensile strength, a layer 0,105 mm thick increased the life of VT14 alloy to 1 hr 15 min, and a layer 0.11 mm thick increased the life of AT8 alloy to 22 hr. The greater strength of the ATS alloy can be explained by its higher aluminum content. Orig. art. has: 3 figures and 1 ASSOCIATION: none SUBMITTED: 00 ENCL: 00 SUB CODE: HM NO REP SOY: 000 OTHER: 000 ATD PRESS: 3150 2/2

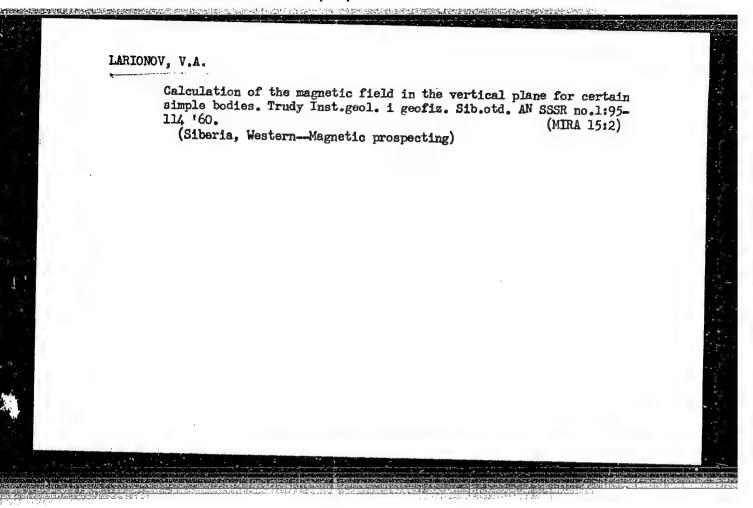
LARIONOV. V.A.

Material and technical security in assembling and in special work. Nov. tekh. i pered. op. v stroi. 20 no. 7:31-32 Jl '58. (MIRA 11:8)

1. Glavnyy inzh. Glavanaba Hinstroya RSFSR.
(Building--Contracts and specifications)



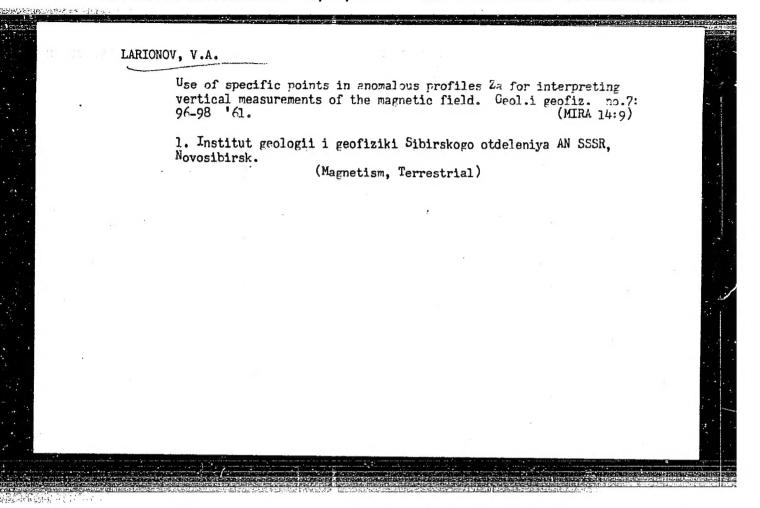
Vertical magnetic logging method. Geol. i geofiz. no.1:107-114 '60. (MEA 13:9) 1. Institut geologii i geofiziki Sibirskogo otdeleniya AN SSSR. (Nagnetic prospecting)

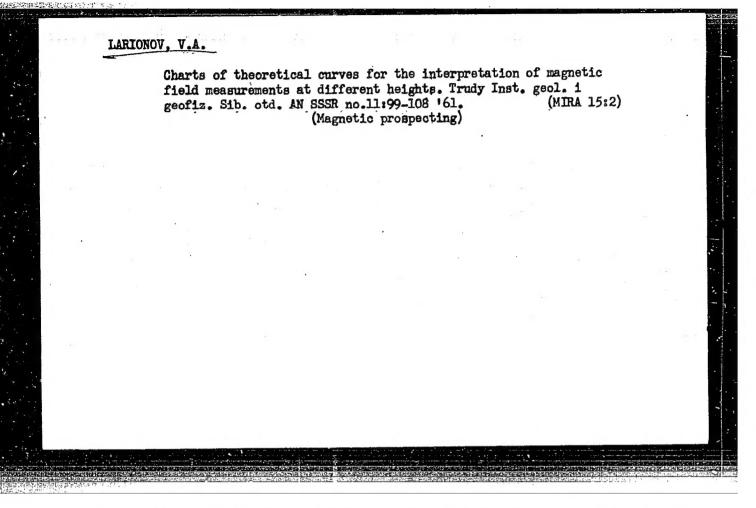


LARIONOV, V. A.

Cand Geol-Min Sci - (diss) "Study of the spatial distribution of magnetic field in surveys and explorations for iron-ore deposits." Novosibirsk, 1961. 19 pp; (Academy of Sciences USSR, Siberian Division, of the Joint Academic Council for Geology-Mineralogy, Geophysics, and Geography); 150 copies; price not given; list of author's works on pp 18-19 (10 entries); (KL, 5-61 sup, 180)

LARIONOV, V.A. Field determination of residual and induced magnetization ratio. Geol. 1 geofiz. no.4:107-109 '61. (MIRA 14:5) 1. Institut geologii i geofiziki Sipirskogo otdeleniya AN SSSR, Novosibirsk. (Magnetism, Terrestrial)





S/169/62/000/007/062/149 D228/D307

AUTHOR:

Larionov, V. A.

TITLE:

Problem of dividing magnetic anomalies into ore and barren ones when seeking and exploring iron ore deposits of the Gornoshorskikh type (Discourse theses)

PERIODICAL:

Card 1/1

Referativnyy zhurnal, Geofizika, no. 7, 1962, 30, abstract 7A196 (V sb. Sostoyaniye i perspektivy razvitiya geofiz. metodov poiskov i razvedki polezn. iskopayemykh, M., Gostoptekhizdat, 1961, 526-527)

TEXT: It was established that the ore complexes of Gornaya Shoriya possess simultaneously heightened magnetic and density properties, the magnetic ores being characterized by the maximum values of the parameters indicated. The execution of vertical plane surveying acquires great significance for determining the positions of the disturbing bodies, with the aim of establishing their ore content. For this case the author developed methods for determining the depth and shape of anomalous masses. / Abstracter's note: Complete translation. /